

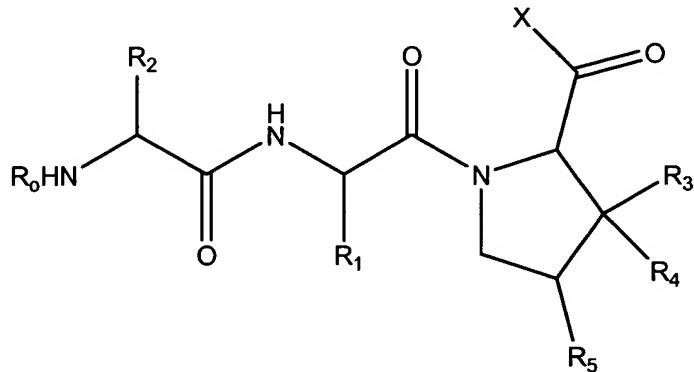


Amendment to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (currently amended): A method for the treatment of a postlesional disease of ischemic, traumatic or toxic origin characterized by nerve cell necrosis, comprising administering an effective amount of a compound of formula (I) to a human patient in need thereof.



(I)

wherein X represents OH, (C₁₋₅)alkoxy, NH₂, NH-(C₁₋₅alkyl), or N(C₁₋₅alkyl)₂ NH-(C₁₋₃)alkyl or N(C₁₋₃alkyl)₂;

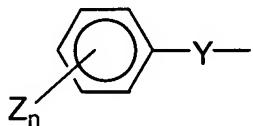
R₁ is a residue derived from any of the amino acid[[s]] Phe, Tyr, Trp, Pro, each of which may optionally be substituted with one or more methoxy groups, or methyl groups by a (C₁₋₅)alkoxy groups, a (C₁₋₅)alkyl group or one or more halogen atoms, and Ala, Val, Leu, ; or is derived from the amino acid Ile;

R₂ is a residue which is derived from any one of the amino acids Gly, Ala, or Ile, Val, Ser, Thr, His, Arg, Lys, Pro, Glu, Gln, pGlu, Asp, Leu or Asn;

R₃-and R₄ independently represent H, OH, (C₁₋₅) alkyl, or (C₁₋₅) alkoxy, provided that R₃ and R₄ are not both OH or (C₁₋₅) alkoxy;

R₅ represents H, OH, (C₁₋₅) alkyl or (C₁₋₅) alkoxy;

and wherein R₀ represents a group of the formula



wherein Y represents -CO-, -CH₂CO-, -CH₂CH₂CO-, CH₂CH₂CH₂CO-, -CH=CH-CO or -OCH₂CO-, and wherein Z represents a halogen atom, a trifluoromethyl group, a methoxy group, a methyl group, (C₁₋₄) alkoxy group, (C₁₋₄) alkyl group; or wherein two neighbouring substituents may form a (C₁₋₃) alkylendioxy group; and wherein n is 0 or an integer of from 1 to 5; or pharmaceutically acceptable salts thereof;

or a pharmaceutically acceptable salt thereof.

2. (Canceled)

3. (Currently Amended) The method according to claim 2, wherein R₁ is a residue derived from Phe which may optionally be substituted by with one or more methoxy groups, or methyl groups a (C₁₋₅) alkoxy groups, a (C₁₋₅) alkyl group or a one or more halogen atoms.

4. (Canceled)

5. (canceled)

6. (previously presented): The method according to claim 1, wherein R₀ is a cinnamoyl moiety.

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7. (previously presented): The method according to claim 1, wherein the compound of formula (I) is cinnamoyl-glycyl-L-phenylalanyl-L-prolinamide, cinnamoyl-isoleucyl-phenylalanyl-L-proline ethylamide, cinnamoyl-isoleucyl-isoleucyl-prolineamide, or a pharmaceutically acceptable salt thereof.